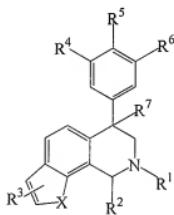


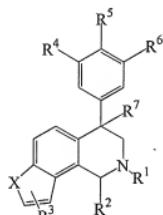
This listing of claims will replace all prior versions, and listings, of claims in the application:

Amendments to the Claims:

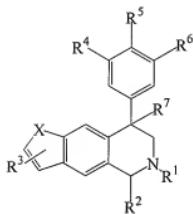
1. (Currently Amended) A method of treating urge-, stress- or mixed-urinary incontinence comprising administration of an effective amount of a compound selected from one of the Formulae IA, IB, IIA, IIB, IIIA or IIIB



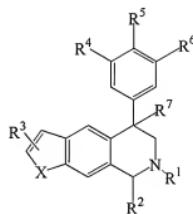
IA



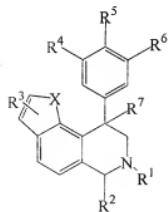
IB



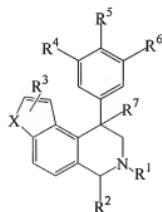
IIA



IIB



IIIA



IIIB

wherein:

PC27831A US Amend & Response DRAFT.doc

R¹ is selected from the group consisting of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₆ cycloalkyl, C₄-C₇ cycloalkylalkyl and benzyl, each of which is optionally substituted with 1 to 3 substituents independently selected at each occurrence from C₁-C₃ alkyl, halogen, -CN, -OR⁸ and -NR⁸R⁹;

R² is selected from the group consisting of H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₆ cycloalkyl, C₄-C₇ cycloalkylalkyl and C₁-C₆ haloalkyl;

R³ is selected from the group consisting of H, halogen, C₁-C₆ alkyl, C₁-C₆ haloalkyl and C₃-C₆ cycloalkyl, wherein C₁-C₆ alkyl, C₁-C₆ haloalkyl and C₃-C₆ cycloalkyl are optionally substituted with 1 to 3 substituents selected independently at each occurrence from OR⁸ and NR⁸R⁹;

R⁴, R⁵, and R⁸ are each independently selected at each occurrence thereof from the group consisting of H, halogen, -OR¹⁰, -NO₂, -NR¹⁰R¹¹, -NR¹⁰C(0)R¹¹, -NR¹⁰C(0)NR¹¹R¹², -S(0)_nR¹¹, -CN, -C(O)R¹¹, -C(O)₂R¹¹, -C(O)NR¹¹R¹², C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₆ cycloalkyl and C₄-C₇ cycloalkylalkyl, wherein each of C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₆ cycloalkyl and C₄-C₇ cycloalkylalkyl are optionally substituted with 1 to 3 substituents independently selected at each occurrence with from C₁-C₃ alkyl, halogen, =O, -CN, -OR⁸, -NR⁸R⁹ and phenyl, and wherein phenyl is optionally substituted 1-3 substituents selected independently at each occurrence from halogen, -CN, C₁-C₄ alkyl, C₁-C₄ haloalkyl, -OR⁸ and -NR⁸R⁹;

alternatively R⁵ and R⁶ taken together are -O-C(R¹¹)₂-O-;

R⁷ is selected from the group consisting of H, halogen and OR¹⁰;

R⁸ and R⁹ are each independently selected from the group consisting of H, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxyalkyl, C₁-C₄ alkoxyalkylalkyl, C₃-C₆ cycloalkyl, C₄-C₇ cycloalkylalkyl, -C(O)R¹², phenyl and benzyl, wherein phenyl and benzyl are optionally substituted with 1 to 3 substituents selected independently at each occurrence from halogen, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy and C₁-C₄ haloalkoxy, or R⁸ and R⁹ are taken together with the nitrogen to which they are attached to form a piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine ring;

R¹⁰ is selected from the group consisting of H, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxyalkyl, C₃-C₆ cycloalkyl, C₄-C₇ cycloalkylalkyl, -C(O)R¹², phenyl and benzyl, wherein phenyl and benzyl are optionally substituted with 1 to 3 substituents selected independently at each

occurrence from halogen, -NH₂, -OH, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy and C₁-C₄ haloalkoxy;

R¹¹ is selected from the group consisting of H, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxyalkyl, C₃-C₆ cycloalkyl, C₄-C₇ cycloalkylalkyl, phenyl and benzyl, where phenyl and benzyl are optionally substituted with 1 to 3 substituents selected independently at each occurrence from halogen, -NH₂, -OH, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy and C₁-C₄ haloalkoxy, or R¹⁰ and R¹¹ are taken together with the nitrogen to which they are attached to form a piperidine, pyrrolidine, N-methylpiperazine, morpholine, or thiomorpholine ring, with the proviso that only one of R⁸ and R9 or R¹⁰ and R¹¹ are taken together with the nitrogen to which they are attached to form a piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine ring;

R¹² is selected from the group consisting of C₁-C₄ alkyl, C₁-C₄ haloalkyl and phenyl;

X is selected from the group consisting of O, NR¹³ and S;

the ring containing X is selected from furan, pyrrole, thiophene, dihydrofuran, dihydropyrrole, and dihydrothiophene; n is 0, 1, or 2; and,

R¹³ is selected from the group consisting of H, C₁-C₆ alkyl, benzyl and phenyl, wherein C₁-C₆ alkyl, benzyl and phenyl are optionally substituted with 1-3 substituents independently at each occurrence from halogen, -NH₂, -OH, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy and C₁-C₄ haloalkoxy;

or a pharmaceutically acceptable salt thereof or an isomer or prodrug thereof to a patient in need thereof.

2. (Original) A method of claim 1, wherein R¹ is C₁-C₆ alkyl.

3. (Original) A method of claim 2, wherein R¹ is CH₃.

4. (Original) A method of claim 1, wherein R² is H, C₁-C₆ alkyl, C₃-C₆ cycloalkyl, or C₁-C₆ haloalkyl.

5. (Original) A method of claim 4, wherein R² is H or C₁-C₆ alkyl.

6. (Original) A method of claim 5, wherein R² is H.

7. (Original) A method of claim 1, wherein R³ is at each occurrence thereof independently H, halogen, C₁-C₆ alkyl, or C₁-C₆ alkyl substituted with from 1 to 3 of OR⁸ or NR⁸R⁹.

8. (Original) A method of claim 7, wherein R³ is H or C₁-C₆ alkyl.

9. (Original) A method of claim 8, wherein R³ is H.

10. (Original) A method of claim 1, wherein R¹ is CH₃, R² is H and R³ is H.

11. (Original) A method of claim 1, wherein R⁴, R⁵ and R⁶ are each independently H, halogen, C₁-C₆ alkyl or -OR¹⁰.

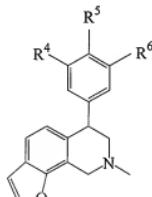
12. (Original) A method of claim 11, wherein at least one of R⁴, R⁵ and R⁶ is H.

13. (Original) A method of claim 12, wherein each of R⁴, R⁵ and R⁶ are H.

14. (Original) A method of claim 12, wherein one of R⁴, R⁵ and R⁶ is halogen.

15. (Original) A method of claim 1, wherein R¹ is CH₃, R² and R³ are each H, and at least one of R⁴, R⁵, and R⁶ is H.

16. (Original) A method of claim 1 wherein the compound is a compound of Formula (10):



(10)

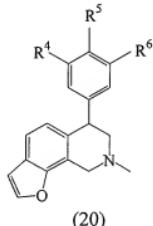
or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (10) wherein R⁴ is H, R⁵ is H and R⁶ is H;

a compound of Formula (10) wherein R⁴ is H, R⁵ is Me and R⁶ is H;

a compound of Formula (10) wherein R⁴ is Cl, R⁵ is H and R⁶ is H; and
 a compound of Formula (10) wherein R⁴ is H, R⁵ is F and R⁶ is H.

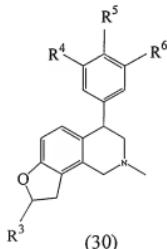
17. (Original) A method of claim 1 wherein the compound is a compound of Formula (20):



or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (20) wherein R⁴ is H, R⁵ is H and R⁶ is H;
 a compound of Formula (20) wherein R⁴ is H, R⁵ is Me and R⁶ is H;
 a compound of Formula (20) wherein R⁴ is H, R⁵ is Cl and R⁶ is H;
 a compound of Formula (20) wherein R⁴ is H, R⁵ is F and R⁶ is H; and
 a compound of Formula (20) wherein R⁴ is F, R⁵ is H and R⁶ is F.

18. (Original) A method of claim 1 wherein the compound is a compound of Formula (30):

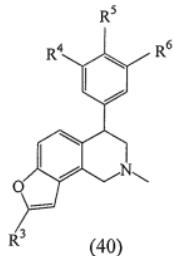


or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (30) wherein R³ is H, R⁴ is H, R⁵ is H and R⁶ is H;
 a compound of Formula (30) wherein R³ is H, R⁴ is F, R⁵ is F and R⁶ is H;
 a compound of Formula (30) wherein R³ is H, R⁴ is F, R⁵ is H and R⁶ is F;

a compound of Formula (30) wherein R³ is H, R⁴ is H, R⁵ is F and R⁶ is H;
 a compound of Formula (30) wherein R³ is H, R⁴ is Cl, R⁵ is H and R⁶ is H;
 a compound of Formula (30) wherein R³ is H, R⁴ is H, R⁵ is Cl and R⁶ is H;
 a compound of Formula (30) wherein R³ is H, R⁴ is H, R⁵ is Cl and R⁶ is F;
 a compound of Formula (30) wherein R³ is H, R⁴ is H, R⁵ is F and R⁶ is Cl;
 a compound of Formula (30) wherein R³ is H, R⁴ is F, R⁵ is H and R⁶ is Cl;
 a compound of Formula (30) wherein R³ is H, R⁴ is F, R⁵ is 0Me and R⁶ is H; and
 a compound of Formula (30) wherein R³ is H, R⁴ is F, R⁵ is H and R⁶ is H.

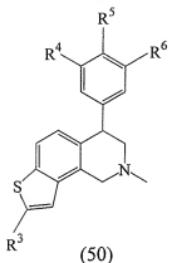
19. (Original) A method of claim 1 wherein the compound is a compound of Formula (40):



or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (40) wherein R³ is H, R⁴ is H, R⁵ is H and R⁶ is H;
 a compound of Formula (40) wherein R³ is H, R⁴ is F, R⁵ is F and R⁶ is H;
 a compound of Formula (40) wherein R³ is H, R⁴ is F, R⁵ is H and R⁶ is F;
 a compound of Formula (40) wherein R³ is H, R⁴ is F, R⁵ is H and R⁶ is H;
 a compound of Formula (40) wherein R³ is H, R⁴ is H, R⁵ is F and R⁶ is H;
 a compound of Formula (40) wherein R³ is H, R⁴ is Cl, R⁵ is H and R⁶ is H;
 a compound of Formula (40) wherein R³ is H, R⁴ is H, R⁵ is Cl and R⁶ is H;
 a compound of Formula (40) wherein R³ is H, R⁴ is H, R⁵ is Cl and R⁶ is F;
 a compound of Formula (40) wherein R³ is H, R⁴ is H, R⁵ is F and R⁶ is Cl;
 a compound of Formula (40) wherein R³ is H, R⁴ is F, R⁵ is H and R⁶ is Cl;
 a compound of Formula (40) wherein R³ is H, R⁴ is H, R⁵ is 0Me and R⁶ is H;
 a compound of Formula (40) wherein R³ is Me, R⁴ is H, R⁵ is H and R⁶ is H;
 a compound of Formula (40) wherein R³ is Et, R⁴ is H, R⁵ is H and R⁶ is H; and
 a compound of Formula (40) wherein R³ is CH₂OH, R⁴ is H, R⁵ is H and R⁶ is H.

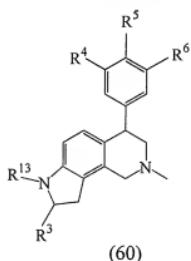
20. (Original) A method of claim 1 wherein the compound is a compound of Formula (50):



or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (50) wherein R³ is H, R⁴ is H, R⁵ is H and R⁶ is H.

21. (Original) A method of claim 1 wherein the compound is a compound of Formula (60):



or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

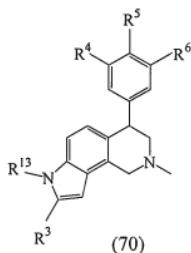
- a compound of Formula (60) wherein R³ is H, R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is H;
- a compound of Formula (60) wherein R³ is H, R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is Me;
- a compound of Formula (60) wherein R³ is H, R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is Et;
- a compound of Formula (60) wherein R³ is H, R⁴ is H, R⁵ is F, R⁶ is F and R¹³ is H;
- a compound of Formula (60) wherein R³ is H, R⁴ is H, R⁵ is F, R⁶ is F and R¹³ is Me;
- a compound of Formula (60) wherein R³ is H, R⁴ is F, R⁵ is H, R⁶ is F and R¹³ is H;
- a compound of Formula (60) wherein R³ is H, R⁴ is F, R⁵ is H, R⁶ is F and R¹³ is Me;

a compound of Formula (60) wherein R³ is H, R⁴ is Cl, R⁵ is H, R⁶ is H and R¹³ is H;
 a compound of Formula (60) wherein R³ is H, R⁴ is Cl, R⁵ is H, R⁶ is H and R¹³ is Me;
 a compound of Formula (60) wherein R³ is H, R⁴ is F, R⁵ is H, R⁶ is H and R¹³ is H;
 a compound of Formula (60) wherein R³ is H, R⁴ is H, R⁵ is F, R⁶ is H and R¹³ is H;
 a compound of Formula (60) wherein R³ is H, R⁴ is F, R⁵ is Cl, R⁶ is H and R¹³ is H;
 a compound of Formula (60) wherein R³ is H, R⁴ is F, R⁵ is Cl, R⁶ is H and R¹³ is Me;
 a compound of Formula (60) wherein R³ is H, R⁴ is Cl, R⁵ is F, R⁶ is H and R¹³ is H;

and

a compound of Formula (60) wherein R³ is H, R⁴ is Cl, R⁵ is F, R⁶ is H and R¹³ is Me.

22. (Original) A method of claim 1 wherein the compound is a compound of Formula (70):



or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

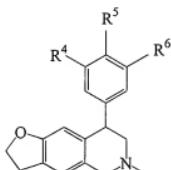
a compound of Formula (70) wherein R³ is H, R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is H;
 a compound of Formula (70) wherein R³ is H, R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is Me;
 a compound of Formula (70) wherein R³ is H, R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is Et;
 a compound of Formula (70) wherein R³ is H, R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is Bn;
 a compound of Formula (70) wherein R³ is H, R⁴ is H, R⁵ is F, R⁶ is F and R¹³ is H;
 a compound of Formula (70) wherein R³ is H, R⁴ is H, R⁵ is F, R⁶ is F and R¹³ is Me;
 a compound of Formula (70) wherein R³ is H, R⁴ is F, R⁵ is H, R⁶ is F and R¹³ is Me;
 a compound of Formula (70) wherein R³ is H, R⁴ is Cl, R⁵ is H, R⁶ is H and R¹³ is H;
 a compound of Formula (70) wherein R³ is H, R⁴ is Cl, R⁵ is H, R⁶ is H and R¹³ is Me;
 a compound of Formula (70) wherein R³ is H, R⁴ is F, R⁵ is H, R⁶ is H and R¹³ is H;
 a compound of Formula (70) wherein R³ is H, R⁴ is F, R⁵ is H, R⁶ is H and R¹³ is Me;
 a compound of Formula (70) wherein R³ is H, R⁴ is F, R⁵ is F, R⁶ is H and R¹³ is H;
 a compound of Formula (70) wherein R³ is H, R⁴ is F, R⁵ is Cl, R⁶ is H and R¹³ is H;

a compound of Formula (70) wherein R³ is H, R⁴ is F, R⁵ is Cl, R⁶ is H and R¹³ is Me;
 a compound of Formula (70) wherein R³ is H, R⁴ is Cl, R⁵ is F, R⁶ is H and R¹³ is H;

and

a compound of Formula (70) wherein R³ is H, R⁴ is Cl, R⁵ is F, R⁶ is H and R¹³ is Me.

23. (Original) A method of claim 1 wherein the compound is a compound of Formula (80):

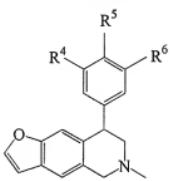


(80)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (80) wherein R⁴ is H, R⁵ is H and R⁶ is H;
 a compound of Formula (80) wherein R⁴ is H, R⁵ is F and R⁶ is H; and
 a compound of Formula (80) wherein R⁴ is H, R⁵ is F and R⁶ is F.

24. (Original) A method of claim 1 wherein the compound is a compound of Formula (90):

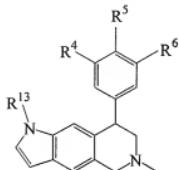


(90)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (90) wherein R⁴ is H, R⁵ is H and R⁶ is H;
 a compound of Formula (90) wherein R⁴ is H, R⁵ is F and R⁶ is F; and
 a compound of Formula (90) wherein R⁴ is H, R⁵ is F and R⁶ is H.

25. (Original) A method of claim 1 wherein the compound is a compound of Formula (100):

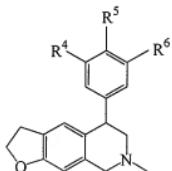


(100)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (100) wherein R^4 is H, R^5 is H, R^6 is H and R^{13} is H.

26. (Currently Amended) A method of claim 1 wherein the compound is a compound of Formula (110):

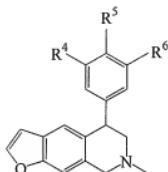


(110)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- a compound of Formula (110) wherein R^4 is H, R^5 is H and R^6 is H;
- a compound of Formula (110) wherein R^4 is H, R^5 is F and R^6 is F;
- a compound of Formula (110) wherein R^4 is H, R^5 is F and R^6 is H;
- a compound of Formula (110) wherein R^4 is H, R^5 is H and R^6 is Cl;
- a compound of Formula (110) wherein R^4 is H, R^5 is Cl and R^6 is F;
- a compound of Formula (110) wherein R^4 is H, R^5 is F and R^6 is Cl; and
- a compound of Formula (110) wherein R^4 is H, R^5 is OMeOMe and R^6 is H.

27. (Currently Amended) A method of claim 1 wherein the compound is a compound of Formula (120):

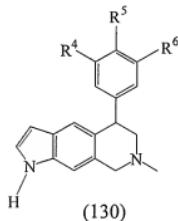


(120)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- a compound of Formula (120) wherein R⁴ is H, R⁵ is H and R⁶ is H;
- a compound of Formula (120) wherein R⁴ is H, R⁵ is F and R⁶ is F;
- a compound of Formula (120) wherein R⁴ is H, R⁵ is F and R⁶ is H;
- a compound of Formula (120) wherein R⁴ is H, R⁵ is H and R⁶ is Cl;
- a compound of Formula (120) wherein R⁴ is H, R⁵ is Cl and R⁶ is F;
- a compound of Formula (120) wherein R⁴ is H, R⁵ is OMe and R⁶ is H; and
- a compound of Formula (120) wherein R⁴ is H, R⁵ is F and R⁶ is Cl.

28. (Original) A method of claim 1 wherein the compound is a compound of Formula (130):

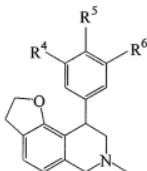


(130)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- a compound of Formula (130) wherein R⁴ is H, R⁵ is H and R⁶ is H; and
- a compound of Formula (130) wherein R⁴ is H, R⁵ is Bn and R⁶ is H.

29. (Currently Amended) A method of claim 1 wherein the compound is a compound of Formula (140):

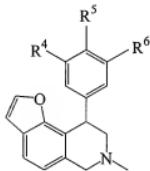


(140)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- a compound of Formula (140) wherein R⁴ is H, R⁵ is H and R⁶ is H;
- a compound of Formula (140) wherein R⁴ is H, R⁵ is F and R⁶ is H;
- a compound of Formula (140) wherein R⁴ is H, R⁵ is F and R⁶ is Cl;
- a compound of Formula (140) wherein R⁴ is H, R⁵ is Cl and R⁶ is F;
- a compound of Formula (140) wherein R⁴ is H, R⁵ is H and R⁶ is Cl;
- a compound of Formula (140) wherein R⁴ is H, R⁵ is OMe and R⁶ is H;
- a compound of Formula (140) wherein R⁴ is H, R⁵ is F and R⁶ is F.

30. (Currently Amended) A method of claim 1 wherein the compound is a compound of Formula (150):



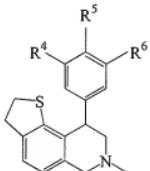
(150)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- a compound of Formula (150) wherein R⁴ is H, R⁵ is H and R⁶ is H;
- a compound of Formula (150) wherein R⁴ is H, R⁵ is F and R⁶ is H;
- a compound of Formula (150) wherein R⁴ is H, R⁵ is F and R⁶ is Cl;
- a compound of Formula (150) wherein R⁴ is H, R⁵ is Cl and R⁶ is F;

- a compound of Formula (150) wherein R⁴ is H, R⁵ is H and R⁶ is Cl;
- a compound of Formula (150) wherein R⁴ is H, R⁵ is OMe and R⁶ is H; and
- a compound of Formula (150) wherein R⁴ is H, R⁵ is F and R⁶ is F.

31. (Original) A method of claim 1 wherein the compound is a compound of Formula (160):

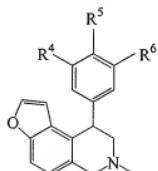


(160)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- a compound of Formula (160) wherein R⁴ is H, R⁵ is H and R⁶ is H.

32. (Original) A method of claim 1 wherein the compound is a compound of Formula (170):

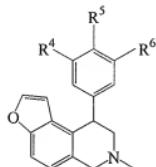


(170)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- a compound of Formula (170) wherein R⁴ is H, R⁵ is H and R⁶ is H;
- a compound of Formula (170) wherein R⁴ is H, R⁵ is F and R⁶ is H; and
- a compound of Formula (170) wherein R⁴ is H, R⁵ is F and R⁶ is F.

33. (Original) A method of claim 1 wherein the compound is a compound of Formula (180):

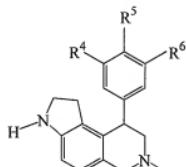


(180)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- a compound of Formula (180) wherein R⁴ is H, R⁵ is H and R⁶ is H;
- a compound of Formula (180) wherein R⁴ is H, R⁵ is F and R⁶ is H; and
- a compound of Formula (180) wherein R⁴ is H, R⁵ is F and R⁶ is F.

34. (Original) A method of claim 1 wherein the compound is a compound of Formula (190):

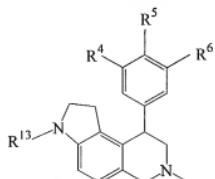


(190)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- a compound of Formula (190) wherein R⁴ is H, R⁵ is H and R⁶ is H.

35. (Original) A method of claim 1 wherein the compound is a compound of Formula (200):



(200)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (200) wherein R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is H; and
a compound of Formula (200) wherein R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is Me.

36. (Currently Amended) A method of claim 1 wherein the compound is selected from the group consisting of:

(R)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2,3-h]isoquinoline;
(S)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2, 3-h]isoquinoline;
(R)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3,2-g]isoquinoline;
(S)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3, 2-g]isoquinoline;
(R)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4- tetrahydrofuro[2,3-h]isoquinoline;
(S)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;
(R)-4-(3,4-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2, 3-h]isoquinoline;
(S)-4-(3, 4-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;
(R)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2, 3-h]isoquinoline;
(S)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2, 3-h]isoquinoline;
(R)-4-(4-chloro-phenyl)-2-methyl-1,2,3,4- tetrahydro-furo[2, 3-h]isoquinoline;
(S)-4-(4-chloro-phenyl)-2- methyl-1,2,3,4-tetrahydrofuro[2,3- h]isoquinoline;
(R)-8-methyl- 6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2-h]isoquinoline;
(S)-8-methyl-6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2-h]isoquinoline;
(R)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;
(S)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;
(R)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3, 4-tetrahydrofuro[2,3-h]isoquinoline;
(S)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(R)-2-methyl-4-phenyl[2,3,4,7-tetrahydro-1H-pyrrolo[2,3-h]isoquinoline; and
(S)-2-methyl-4-phenyl-2,3,4,7-tetrahydro-1H-pyrrolo[2, 3-h]isoquinoline; or a pharmaceutically acceptable salts thereof.

37. (Currently Amended) A method of claim 1 wherein the compound is selected from the group consisting of:

(+)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2, 3-h]isoquinoline;
(-)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2,3-h]isoquinoline;
(+)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3, 2-g]isoquinoline;
(-)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3, 2-g]isoquinoline;

(+)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(-)4-(4-fluoro-phenyl)-2-methyl-1,2,3,4- tetrahydro-furo[2,3-h]isoquinoline;
(+)-4-(3,4-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(-)4-(3,4- difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(+)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(-)2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(+)-4-(4-chloro-phenyl)-2-methyl-1,2,3,4- tetrahydrofuro[2,3-h]isoquinoline;
(-)4-(4-chloro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(+)-8-methyl-6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2h]isoquinoline;
(-)8-methyl- 6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2-h]isoquinoline;
(+)-4-(4- fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(-)4-(4-fluoro-phenyl)-2-methyl-1,2,3,4- tetrahydrofuro[2,3-h]isoquinoline;
(+)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(-)4-(3,5-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
(+)-2-methyl-4-phenyl-2,3,4,7-tetrahydro-1H-pyrrolo[2,3-h]isoquinoline; and
(-)2-methyl-4-phenyl-2,3,4,7-tetrahydro-1H- pyrrolo[2,3-h]isoquinoline; or a
pharmaceutically acceptable salts thereof.